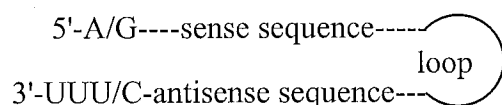


IN THE CLAIMS

The claims are as follows:

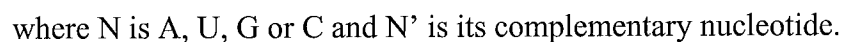
1. (Original) A recombinant vector for the correct, stable and effective expression in mammalian cells of a siRNA or a miRNA, comprising from 5' to 3':
 - a) an RNA polymerase II dependent promoter sequence derived from the U1 snRNA gene;
 - b) suitable restriction sites for cloning the sequence that transcribes a presiRNA or a pre-miRNA;
 - c) a sequence transcribing the pre-siRNA comprising: in position +1 an A or a G residue; a sequence from 21 to 23 nucleotides corresponding to a sense region of the mRNA transcribed by the gene to be silenced, that constitutes the first segment of the stem of the pre-siRNA; a sequence selected from a pre-miRNA sequence that constitutes the loop region of the pre-siRNA; a sequence from 21 to 23 nucleotides corresponding to the antisense region of the mRNA transcribed by the gene to be silenced that constitutes the second segment of the stem of the pre-siRNA; two final residues UU protruding in such a way that the following structure is obtained:



or alternatively a sequence transcribing the pre-miRNA;

- d) termination sequences derived from the sequence at 3' of the gene for U1 snRNA which are necessary and sufficient for the correct formation of the 3' of the pre-siRNA or of the pre-miRNA.

2. (Original) The vector according to Claim 1, wherein the cloning site for the 5' of the sequence transcribing the pre-siRNA is Bgl II.
3. (Original) The vector according to Claim 1, wherein the sequence transcribing the pre-siRNA further comprises at termini 5' and 3' such sequence that the transcribed pre-siRNA has the following structure:



5'-AUA A----sense sequence-
 3'-GUCCCCUAU U--antisense sequence--
 (SEQ ID NO:1)

CCCCTG/ACTTTCTGGAGTTTCAAAGTAGAC (SEQ ID NO:18).

8. (Canceled).